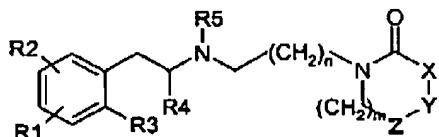


Atty Docket No.: R0067D DIV
USSN: 10/685,124

CLAIM LISTING

Claims 1-41 (canceled)

42. (Currently Amended) A compound of Formula I



wherein:

R¹, R² and R³ are independently in each occurrence hydrogen, halogen, (C₁₋₆) - alkyl, -OR', -SR', -NR'R'', -SOR', -SO₂R', -COOR', -OCOR', -OCONRR'', -OSO₂R', -OSO₂NR'R''; -NR'SO₂R'', -NR'COR'', -SO₂NR'R'', -SO₂(CH₂)₁₋₃CONR'R'', -CONRR'', cyano, haloalkyl, or nitro; or R¹ and R² if adjacent, taken together with the carbons to which they are attached may also form a 5- to 7- membered aromatic, saturated or unsaturated ring, optionally incorporating one or two ring heteroatoms chosen from N, S (O)₀₋₂, or O, and optionally substituted with (C₁₋₆)-alkyl, halo, cyano or lower alkoxy;

R' and R'' are independently in each occurrence hydrogen, (C₁₋₆)-alkyl, substituted lower alkyl, (C₀₋₃)alkylalkoxy, aryl, heterocyclyl, heteroaryl, aryl-(C₁₋₃)-alkyl, heteroaryl-(C₁₋₃)-alkyl, heterocyclyl-(C₁₋₃)-alkyl, cycloalkylalkyl, cycloalkyl, or R' and R'' together with the nitrogen they are attached may also form a 5- to 7- membered ring, optionally incorporating one additional ring heteroatom chosen from N, O or S(O)₀₋₂;

R⁴ is independently in each occurrence (C₁₋₆) alkyl;

R⁵ is independently in each occurrence (C₁₋₆) alkyl, (C₁₋₆) alkenyl, (C₁₋₆) alkynyl, or cycloalkyl;

one of X, Y or Z is independently S, O, or N-R⁶, the others are CH₂;

R⁶ is hydrogen, (C₁₋₆)-alkyl, haloalkyl, aryl(C₁₋₆)alkyl, heteroaryl(C₁₋₆)alkyl, -(C₁₋₆)-CR'R'R'', -COOR', -SO₂R', -C(O)R', -SO₂(CH₂)₀₋₃NR'R'', -CONR'R'', -C(O)OCH₂OC(O)R', -C(O)OCH₂SC(O)R', or -PO(OR')₂, where R' and R'' are as defined above;

m is 1;

Atty Docket No.: R0067D DIV
USSN: 10/685,124

n is 3 ~~an integer from 1 to 6 inclusive;~~
or pharmaceutically acceptable salts or solvates thereof.

43. (Canceled)
44. (Previously Presented) The compound of Claim 42, wherein R⁴ is methyl.
45. (Canceled)
46. (Previously Presented) The compound of Claim 42, wherein X is S or O.
47. (Previously Presented) The compound of Claim 42, wherein Y is S or O.
48. (Previously Presented) The compound of Claim 42, wherein Z is S or O.
49. (Previously Presented) The compound of Claim 42, wherein one of X, Y or Z is NR⁶ and the others are CH₂.
50. (Previously Presented) The compound of Claim 49, wherein X is NH.
51. (Previously Presented) The compound of Claim 49, wherein Y is NH.
52. (Previously Presented) The compound of Claim 49, wherein Z is NH.
53. (Previously Presented) The compound of claim 42, wherein X is S, O, or N-R⁶, and Y and Z are CH₂.
54. (Currently Amended) The compound of claim 41[3]12, wherein X is S or O, and Y and Z are CH₂.

Atty Docket No.: R0067D DIV
USSN: 10/685,124

55. (Previously Presented) A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 42 in admixture with a pharmaceutically acceptable carrier.

56. (Canceled)

57. (Previously Presented) A method for treating a subject suffering from detrusor hyperactivity ~~a smooth muscle function disease~~ mediated by an M2/M3 muscarinic receptor antagonist, said method comprising administering to said subject an effective amount of at least one compound of claim 42.

58. (Canceled)

59. (Canceled)